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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
09/810,796	03/15/2001	Timothy J. Jegla	018512-005010US	6783

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EXAMINER

BUNNER, BRIDGET E

ART UNIT PAPER NUMBER

1647

DATE MAILED: 04/30/2002

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Please find below and/or attached an Office communication concerning this application or proceeding.

Office Action Summary

Application No.

09/810,796

Applicant(s)

JEGLA, TIMOTHY J.

Examiner

Bridget E. Bunner

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-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --

Period for Reply

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 1 MONTH(S) FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If the period for reply specified above is less than thirty (30) days, a reply within the statutory minimum of thirty (30) days will be considered timely.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133).
- Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

Status

- 1) ☒ Responsive to communication(s) filed on 01 February 2002.
- 2a) ☐ This action is **FINAL**. 2b) ☒ This action is non-final.
- 3) ☐ Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

Disposition of Claims

- 4) ☒ Claim(s) 1-40 is/are pending in the application.
- 4a) Of the above claim(s) _____ is/are withdrawn from consideration.
- 5) ☐ Claim(s) _____ is/are allowed.
- 6) ☐ Claim(s) _____ is/are rejected.
- 7) ☐ Claim(s) _____ is/are objected to.
- 8) ☒ Claim(s) 1-40 are subject to restriction and/or election requirement.

Application Papers

- 9) ☐ The specification is objected to by the Examiner.
- 10) ☐ The drawing(s) filed on _____ is/are: a) ☐ accepted or b) ☐ objected to by the Examiner.
- Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).
- 11) ☐ The proposed drawing correction filed on _____ is: a) ☐ approved b) ☐ disapproved by the Examiner.
- If approved, corrected drawings are required in reply to this Office action.
- 12) ☐ The oath or declaration is objected to by the Examiner.

Priority under 35 U.S.C. §§ 119 and 120

- 13) ☐ Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
- a) ☐ All b) ☐ Some * c) ☐ None of:
- ☐ Certified copies of the priority documents have been received.
 - ☐ Certified copies of the priority documents have been received in Application No. _____.
 - ☐ Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).
- * See the attached detailed Office action for a list of the certified copies not received.
- 14) ☐ Acknowledgment is made of a claim for domestic priority under 35 U.S.C. § 119(e) (to a provisional application).
- a) ☐ The translation of the foreign language provisional application has been received.
- 15) ☐ Acknowledgment is made of a claim for domestic priority under 35 U.S.C. §§ 120 and/or 121.

Attachment(s)

- ☐ Notice of References Cited (PTO-892)
- ☐ Notice of Draftsperson's Patent Drawing Review (PTO-948)
- ☐ Information Disclosure Statement(s) (PTO-1449) Paper No(s) _____
- ☐ Interview Summary (PTO-413) Paper No(s) _____
- ☐ Notice of Informal Patent Application (PTO-152)
- ☐ Other:

DETAILED ACTION

Election/Restrictions

1. Restriction to one of the following inventions is required under 35 U.S.C. 121:
 - A. Claims 1-11, 22-23, drawn to an isolated nucleic acid, an expression vector, and a host cell, classified in class 536, subclass 23.1.
 - B. Claim 12, drawn to a method of detecting a nucleic acid, classified in class 435, subclass 6.
 - C. Claims 13-19, drawn to an isolated polypeptide comprising an alpha subunit of KCNQ potassium channel, classified in class 530, subclass 350.
 - D. Claims 20-21, drawn to an antibody that specifically binds to the KCNQ polypeptide, classified in class 530, subclass 387.1.
 - E. Claims 24-34, drawn to a method for identifying a compound that increases or decreases ion flux through a potassium channel, classified in class 435, subclass 7.1.
 - F. Claim 35, drawn to a method for modulating ion flux through a KCNQ potassium channel, classified in class 435, subclass 7.1.
 - G. Claim 36, drawn to a method for identifying a compound that increases or decreases ion flux through a potassium channel comprising a KCNQ5 polypeptide, classified in class 435, subclass 4.
 - H. Claims 37-38, drawn to a method of detecting the presence of hKCNQ5 in human tissue, classified in class 435, subclass 7.1.
 - I. Claims 39-40, drawn to a method of screening for mutations of a human KCNQ5 gene, classified in class 435, subclass 4.

The inventions are distinct, each from the other because of the following reasons:

- a. Although there are no provisions under the section for "Relationship of Inventions" in M.P.E.P. § 806.05 for inventive groups that are directed to different products, restriction is deemed to be proper because these products constitute patentably distinct inventions for the following reasons. Groups A, C-

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D are directed to products that are distinct both physically and functionally, are not required one for the other, and are therefore patentably distinct. Further, the protein of Group C can be prepared by processes which are materially different from recombinant DNA expression of Group A, such as by chemical synthesis, or by isolation and purification from natural sources. Additionally, the DNA of Group A can be used other than to make the protein of Group C, such in gene therapy or as a probe in nucleic acid hybridization assays. The protein of Group C can be used in materially different methods other than to make the antibody of Group D, such as in therapeutic or diagnostic methods (e.g., in screening). Finally, although the antibody of Group D can be used to obtain the DNA of Group A, it can also be used in materially different methods, such as in various diagnostic (e.g., as a probe in immunoassays or immunochromatography), or therapeutic methods.

- b. Similarly, although there are no provisions under the section for "Relationship of Inventions" in M.P.E.P. § 806.05 for inventive groups that are directed to different methods, restriction is deemed to be proper because these methods constitute patentably distinct inventions for the following reasons. Inventions B and E-I are different methods because they require different ingredients, process steps, and endpoints. Groups B and E-I are different methods requiring different method steps, wherein each is not required, one for another. For example, Invention B requires search and consideration of detection of a nucleic acid by contacting the nucleic acid with an isolated nucleic acid, which is not required by the other inventions. Invention E requires search and consideration of identification of a compound that increases or decreases ion flux through a potassium channel by contacting the compound with a KCNQ polypeptide and determining the functional effect of the compound upon the potassium channel, which is not required by the other inventions. Invention F requires search and consideration of modulating ion flux through a potassium channel by contacting the KCNQ potassium channel with an identified compound, which is not required

by the other inventions. Invention G requires search and consideration of identification of a compound that increases or decreases ion flux through a potassium channel by entering into a computer system an amino acid sequence or nucleotide sequence of KCNQ5, generating a 3-D structure of the polypeptide, generating a 3-D structure of the potassium channel comprising the KCNQ5 polypeptide, generating a 3-D structure of the compound, and comparing the 3-D structures of the polypeptide and compound, which is not required by the other inventions. Invention H requires search and consideration of detection of hKCNQ5 in human tissue by isolating a biological sample, contacting the sample with a KCNQ5-specific reagent, and detecting the level of hKCNQ5-specific reagent that selectively associates with the sample, which is not required by the other inventions. Invention I requires search and consideration of screening for mutations of human KCNQ5 gene by using a computer system, which is not required by the other inventions.

- c. Inventions A and B/I are related as product and process of use. The inventions can be shown to be distinct if either or both of the following can be shown: (1) the process for using the product as claimed can be practiced with another materially different product or (2) the product as claimed can be used in a materially different process of using that product (MPEP § 806.05(h)). In the instant case, the product claimed can be used in materially different processes, such as DNA purification and gene therapy.
- d. Inventions C and E/F/G are related as product and process of use. The inventions can be shown to be distinct if either or both of the following can be shown: (1) the process for using the product as claimed can be practiced with another materially different product or (2) the product as claimed can be used in a materially different process of using that product (MPEP § 806.05(h)). In the instant case, the product claimed can be used in materially different assays, such as an antigen for the production of antibodies.

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- e. Inventions A and E/F/G/H are unrelated. Inventions are unrelated if it can be shown that they are not disclosed as capable of use together and they have different modes of operation, different functions, or different effects (MPEP § 806.04, MPEP § 808.01). In the instant case, the different inventions of Groups A and E/F/G/H are unrelated products and methods, wherein each is not required, one for another. For example, the nucleic acid of Invention A cannot be used together with the claimed methods of Inventions E/F/G/H because these inventions do not recite the use or production of the nucleic acid of Invention A.
 - f. Inventions C and B/H/I are unrelated. Inventions are unrelated if it can be shown that they are not disclosed as capable of use together and they have different modes of operation, different functions, or different effects (MPEP § 806.04, MPEP § 808.01). In the instant case, the different inventions of Groups C and B/H/I are unrelated products and methods, wherein each is not required, one for another. For example, the polypeptide of Invention C cannot be used together with the claimed methods of Inventions B/H/I because these inventions do not recite the use or production of the polypeptide of Invention C.
 - g. Inventions D and B/E/F/G/H/I are unrelated. Inventions are unrelated if it can be shown that they are not disclosed as capable of use together and they have different modes of operation, different functions, or different effects (MPEP § 806.04, MPEP § 808.01). In the instant case, the different inventions of Groups D and B/E/F/G/H/I are unrelated products and methods, wherein each is not required, one for another. For example, the antibody of Invention D cannot be used together with the claimed methods of Inventions B/E/F/G/H/I because these inventions do not recite the use or production of the antibody of Invention D.
2. Because these inventions are distinct for the reasons given above and have acquired a separate status in the art because of their separate search requirements, different classification,

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and recognized divergent subject matter, restriction for examination purposes as indicated is proper.

3. Restriction to one of the following inventions is also required under 35 U.S.C. 121:

Groups 1-3. The inventions as they pertain to each of SEQ ID NOS: 1-3, classification dependent upon the nature of the inventions.

The inventions are distinct, each from the other because of the following reasons:

- h. Although there are no provisions under the section for "Relationship of Inventions" in M.P.E.P. § 806.05 for inventive groups that are directed to different products, restriction is deemed to be proper because these products constitute patentably distinct inventions for the following reasons. Each of SEQ ID NOS: 1-3 is a unique nucleic acid sequence, requiring a unique search of the prior art. Searching all of the sequences in a single patent application would provide an undue search burden on the examiner and the USPTO's resources because of the non-coextensive nature of these searches.

Because these inventions are distinct for the reasons given above and have acquired a separate status in the art as shown by their separate search requirements and recognized divergent subject matter, restriction for examination purposes as indicated is proper.

4. Restriction to one of the following inventions is also required under 35 U.S.C. 121:

Groups 4-5. The inventions as they pertain to each of SEQ ID NOS: 4-5, classification dependent upon the nature of the inventions.

The inventions are distinct, each from the other because of the following reasons:

- i. Although there are no provisions under the section for "Relationship of Inventions" in M.P.E.P. § 806.05 for inventive groups that are directed to different products, restriction is deemed to be proper because these products constitute patentably distinct inventions for the following reasons. Each of SEQ

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ID NOS: 4-5 is a unique amino acid sequence, requiring a unique search of the prior art. Searching all of the sequences in a single patent application would provide an undue search burden on the examiner and the USPTO's resources because of the non-coextensive nature of these searches.

Because these inventions are distinct for the reasons given above and have acquired a separate status in the art as shown by their separate search requirements and recognized divergent subject matter, restriction for examination purposes as indicated is proper.

5. This application contains claims directed to the following patentably distinct species of the claimed invention:

The nucleic acid amplified by primers that selectively hybridize under stringent conditions to the same sequence of primers selected from:

- aa. SEQ ID NO: 6
- bb. SEQ ID NO: 7
- cc. SEQ ID NO: 8
- dd. SEQ ID NO: 9
- ee. SEQ ID NO: 10
- ff. SEQ ID NO: 11
- gg. SEQ ID NO: 12
- hh. SEQ ID NO: 13

Applicant is required under 35 U.S.C. 121 to elect a single disclosed species for prosecution on the merits to which the claims shall be restricted if no generic claim is finally held to be allowable. Currently, claims 2-5 and 7-40 are generic.

Applicant is advised that a reply to this requirement must include an identification of the species that is elected consonant with this requirement, and a listing of all claims readable thereon, including any claims subsequently added. An argument that a claim is allowable or that all claims are generic is considered nonresponsive unless accompanied by an election.

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Upon the allowance of a generic claim, applicant will be entitled to consideration of claims to additional species which are written in dependent form or otherwise include all the limitations of an allowed generic claim as provided by 37 CFR 1.141. If claims are added after the election, applicant must indicate which are readable upon the elected species. MPEP § 809.02(a).

Should applicant traverse on the ground that the species are not patentably distinct, applicant should submit evidence or identify such evidence now of record showing the species to be obvious variants or clearly admit on the record that this is the case. In either instance, if the examiner finds one of the inventions unpatentable over the prior art, the evidence or admission may be used in a rejection under 35 U.S.C. 103(a) of the other invention.

6. This application contains claims directed to the following patentably distinct species of the claimed invention:

A method of identifying a compound that increases or decreases ion flux through a potassium channel by contacting the compound with a KCNQ polypeptide and determining the functional effect of the compound wherein the functional effect is determined by measuring:

- ii. ion flux
- jj. changes in ion concentrations
- kk. changes in current
- ll. changes in voltage
- mm. ligand binding to the channel

Applicant is required under 35 U.S.C. 121 to elect a single disclosed species for prosecution on the merits to which the claims shall be restricted if no generic claim is finally held to be allowable. Currently, claims 1-23, 25-26, and 30-40 are generic.

Applicant is advised that a reply to this requirement must include an identification of the species that is elected consonant with this requirement, and a listing of all claims readable thereon, including any claims subsequently added. An argument that a claim is allowable or that all claims are generic is considered nonresponsive unless accompanied by an election.

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Upon the allowance of a generic claim, applicant will be entitled to consideration of claims to additional species which are written in dependent form or otherwise include all the limitations of an allowed generic claim as provided by 37 CFR 1.141. If claims are added after the election, applicant must indicate which are readable upon the elected species. MPEP § 809.02(a).

Should applicant traverse on the ground that the species are not patentably distinct, applicant should submit evidence or identify such evidence now of record showing the species to be obvious variants or clearly admit on the record that this is the case. In either instance, if the examiner finds one of the inventions unpatentable over the prior art, the evidence or admission may be used in a rejection under 35 U.S.C. 103(a) of the other invention.

7. This application contains claims directed to the following patentably distinct species of the claimed invention:

A method of detecting the presence of hKCNQ5 in human tissue by isolating a biological sample and contacting the biological sample with a hKCNQ5-specific reagent wherein the hKCNQ6-specific reagent is:

- nn. hKCNQ5-specific antibodies
- oo. hKCNQ5-specific oligonucleotide primers
- pp. hKCNQ5-specific nucleic acid probes

Applicant is required under 35 U.S.C. 121 to elect a single disclosed species for prosecution on the merits to which the claims shall be restricted if no generic claim is finally held to be allowable. Currently, claims 1-36 and 39-40 are generic.

Applicant is advised that a reply to this requirement must include an identification of the species that is elected consonant with this requirement, and a listing of all claims readable thereon, including any claims subsequently added. An argument that a claim is allowable or that all claims are generic is considered nonresponsive unless accompanied by an election.

Upon the allowance of a generic claim, applicant will be entitled to consideration of claims to additional species which are written in dependent form or otherwise include all the limitations of an allowed generic claim as provided by 37 CFR 1.141. If claims are added after

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the election, applicant must indicate which are readable upon the elected species. MPEP § 809.02(a).

Should applicant traverse on the ground that the species are not patentably distinct, applicant should submit evidence or identify such evidence now of record showing the species to be obvious variants or clearly admit on the record that this is the case. In either instance, if the examiner finds one of the inventions unpatentable over the prior art, the evidence or admission may be used in a rejection under 35 U.S.C. 103(a) of the other invention.

Applicant is advised that the reply to this requirement to be complete must include an election of the invention to be examined even though the requirement be traversed (37 CFR 1.143).

In order to be fully responsive, Applicant must select one from Groups A-I, one from 1-3 and one from 4-5. Applicant is advised that A-I, 1-3, and 4-5 are not species election requirements; rather, each of A-I, 1-3, and 4-5 is a restriction requirement.

If Applicant selects invention A, one species from the primer sequence group must be chosen to be fully responsive.

If Applicant selects invention E, one species from the determination of functional effect group must be chosen to be fully responsive.

If Applicant selects invention H, one species from the hKCNQ5-specific reagent group must be chosen to be fully responsive.

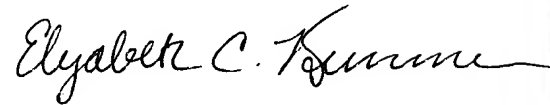
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Any inquiry concerning this communication or earlier communications from the examiner should be directed to Bridget E. Bunner whose telephone number is (703) 305-7148. The examiner can normally be reached on 8:00-5:30 M-F.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Gary Kunz can be reached on (703) 308-4623. The fax phone numbers for the organization where this application or proceeding is assigned are (703) 308-4242 for regular communications and (703) 308-4242 for After Final communications.

Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the receptionist whose telephone number is (703) 308-0196.

BEB
Art Unit 1647
March 28, 2002



ELIZABETH KEMMERER
PRIMARY EXAMINER